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ABSTRACT

INDOLE-AMIDE DERIVATIVES AND THEIR USE AS GLYCOGEN PHOSPHORYLASE INHIBITORS

Heterocyclic amides of formula (1)

wherein:

R¹ is independently selected from, for example, C₁₋₆alkyl, C₅₋₇cycloalkyl,

 C_{5-7} cycloalkyl C_{1-3} alkyl, C_{1-6} alkoxy, C_{5-7} cycloalkoxy, C_{5-7} cycloalkyl C_{1-3} alkoxy, heterocyclyl, heterocyclyl C_{1-3} alkyl, heterocyclyloxy or heterocyclyl C_{1-3} alkoxy;

R² is phenyl or heteroaryl;

R³ is independently selected from hydrogen, halo, nitro, cyano, hydroxy, carboxy, carbamoyl, C₁₋₄alkyl, C₂₋₄alkenyl, C₂₋₄alkynyl, C₁₋₄alkoxy, C₁₋₄alkanoyl, fluoromethyl, difluoromethyl, trifluoromethyl and trifluoromethoxy;

or a pharmaceutically acceptable salt or pro-drug thereof; possess glycogen phosphorylase inhibitory activity and accordingly have value in the treatment of disease states associated with increased glycogen phosphorylase activity. Processes for the manufacture of said heterocyclic amide derivatives and pharmaceutical compositions containing them are described.